AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application:

Listing of Claims:

1. (Currently Amended) A compound of formula I

wherein

 $X \text{ is } = CR^0 - \text{ or } = N -;$

- each of R⁰, R¹, R², R³ and R⁴ independently is hydrogen; hydroxy; C₁-C₈alkyl; C₂-C₈alkenyl; C₃-C₈cycloalkyl; C₃-C₈cycloalkyl-C₁-C₈alkyl; hydroxyC₁-C₈alkyl; C₁-C₈alkoxyC₁-C₈alkyl; hydroxyC₁-C₈alkoxyC₁-C₈alkyl; arylC₁-C₈alkyl which optionally may be substituted on the ring by hydroxy, C₁-C₈alkoxy, carboxy or C₁-C₈alkoxycarbonyl;
- or R³ and R⁴ form together with the nitrogen and carbon atoms to which they are attached a 5 to 10 membered heterocyclic ring and having 1, 2 or 3 heteroatoms selected from N, O and S;
- or each of R¹, and R² and R², independently, is halogen; halo-C₁-C₈alkyl; C₁-C₈alkoxy; halo-C₁-C₈alkoxy; hydroxyC₁-C₈alkoxy; C₁-C₈alkoxyC₁-C₈alkoxy; aryl; arylC₁-C₈alkoxy; heteroaryl; heteroaryl-C₁-C₄alkyl; 5 to 10 membered heterocyclic ring; nitro; carboxy;
- C_2 - C_8 alkoxycarbonyl; C_2 - C_8 alkylcarbonyl; -N(C_1 - C_8 alkyl)C(O) C_1 - C_8 alkyl; -N(R^{10}) R^{11} ;
- -CON(R^{10}) R^{11} ; -SO₂N(R^{10}) R^{11} ; or -C₁-C₄-alkylene-SO₂N(R^{10}) R^{11} ; wherein each of R^{10} and R^{11} independently is hydrogen; hydroxy; C₁-C₈alkyl; C₂-C₈alkenyl; C₃-C₈cycloalkyl;
- C₃-C₈cycloalkyl-C₁-C₈alkyl; C₁-C₈alkycyC₁-C₈alkyl; hydroxyC₁-C₈alkycyC₁-C₈alkyl; hydroxyC₁-C₈alkyl)-carbonyl; arylC₁-C₈alkyl which

optionally may be substituted on the ring by hydroxy, C₁-C₈alkoxy, carboxy or C₂-C₈alkoxycarbonyl; or 5 to 10 membered heterocyclic ring;

or R¹ and R² form together with the C-atoms to which they are attached aryl or a 5 to 10 membered heteroaryl group having one or two heteroatoms selected from N, O and S; or

R⁵ is hydrogen; halogen; cyano; C₁-C₈alkyl; halo-C₁-C₈alkyl;

 C_2 - C_8 alkenyl; C_2 - C_8 alkynyl; C_3 - C_8 cycloalkyl; C_3 - C_8 cycloalkyl; C_5 - C_{10} aryl C_1 - C_8 alkyl;

R⁶ is hydrogen;

each of R⁷, R⁸ and R⁹ is independently hydrogen; hydroxy; C₁-C₈alkyl; C₂-C₈alkenyl; halo-C₁-C₈alkyl; C₁-C₈alkoxy; C₃-C₈cycloalkyl; C₃-C₈cycloalkylC₁-C₈alkyl; arylC₁-C₈alkyl; -Y-R¹² wherein Y is a direct bond or O and R¹² is a substituted or unsubstituted 5, 6 or 7 membered heterocyclic ring comprising 1, 2 or 3 heteroatoms selected from N, O and S; carboxy; (C₁-C₈alkoxy)-carbonyl; -N(C₁₋₈alkyl)-CO-NR¹⁰R¹¹; -CONR¹⁰R¹¹; -N(R¹⁰)(R¹¹); or R⁷ and R⁸ or R⁸ and R⁹, respectively form together with the carbon atoms to which they are attached, a 5 or 6 membered heteroaryl comprising 1, 2 or 3 heteroatoms selected from N, O and S; or a 5 or 6 membered carbocyclic ring;

provided that one of R^4 , R^2 or R^3 is $-CON(R^{10})R^{14}$ or $-SO_2N(R^{10})R^{11}$; in free form or salt form;

wherein

aryl represents phenyl, naphthyl or 1,2,3,4-tetrahydronaphthyl,

heteroaryl is a 5 or 6 membered aromatic heterocyclic ring, optionally condensed to 1 or 2 benzene rings and/or to a further heterocylic ring, and

wherein a heterocyclic ring is a 5 or 6 membered heterocyclic ring being saturated or unsaturated and optionally condensed to 1 or 2 benzene rings and/or to a further heterocyclic ring

2. (Original) A process for the production of a compound of formula I according to claim 1, comprising the steps of reacting a compound of formula II wherein R¹, R², R³, R⁴, R⁵, R⁶ and X are as defined in claim 1, and Y is a leaving group; with a compound of formula III wherein R⁷, R.sup.8 and R.sup.9 are as defined in

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claim 1; and recovering the resulting compound of formula I in free form or in salt form, and, where required, converting the compound of formula I obtained in free form into the desired salt form, or vice versa.

- 3. (Cancelled).
- 4. (Original) A pharmaceutical composition comprising a compound of formula I according to claim 1 or a pharmaceutically acceptable salt thereof, together with one or more pharmaceutically acceptable carriers or diluents therefor.
- 5-7. (Cancelled).
- 8. (Previously Presented) A method for treating breast cancer, comprising: administering to a subject in need thereof, a therapeutically effective amount of a compound of formula I according to claim 1 or a pharmaceutically acceptable salt thereof.
- 9. (Cancelled).